REMARKS

The present response is submitted in reply to the Final Office Action issued on January 8, 2009. Claims 1-5, 7, 8 and 10 are pending in this application, all of which have been rejected. By the present response, claims 1 and 3 have been amended. No new matter has been added.

Reconsideration is respectfully requested in light of the following remarks.

Rejection of claims 1-5, 7, 8 and 10 under 35 U.S.C. 112, second paragraph

Claims 1-5, 7, 8 and 10 have been rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which the applicants regard as the invention. The Examiner states that it is still not clear what is meant by the phrase "the residue of an amino acid" and asks whether the fragment includes amine, carboxylic acid, esters, alcohols or even hydrogen.

It is submitted that the aforementioned phrase has been clarified, as set forth above in the amended claims. In particular, the phrase has been amended to recited "alpha amino acid" and to define the "reside of an amino acid" by adding "the N of the ring is acylated by the carboxylic group of an amino acid" with respect to R_5 and R_7 . Support can be found in the specification, such as at page 6, first paragraph.

The Applicants further submit that "the residue of an amino acid" should nevertheless be clear to one skilled in the art and that in no case can be hydrogen. One skilled in the art should readily recognize what such a residue would be, i.e., that the residue may be the residue of a naturally occurring alpha amino acid as set forth in the specification (page 6, first paragraph). The Applicants would be glad to provide a list of the naturally occurring alpha amino acids at the Examiner's request. In this regard, it is

worth pointing out that the term "residue of an amino acid" was considered definite in U.S. Patent No. 6,753,445 (e.g., claim 3), the patent having at least one inventor in common with the present application.

In view of the above, withdrawal of the rejection is requested.

Rejection of claims 1-10 under 35 U.S.C. 103(a)

Claims 1-5, 7, 8 and 10 are rejected under 35 U.S.C. 103(a) as being unpatentable over U.S. Patent No. 7,169,804 (Ascher, et al.) and/or U.S. Patent No. RE39128 (Berry, et al.). In particular, the Examiner argues that Ascher, et al. and Berry, et al. teach antibacterial mutilin compounds and compositions corresponding to those recited in the present claims, and refers to column 1, line 1 through column 2, line 34 of Ascher, et al. and column 2, lines 2-52, column 3, lines 5-10 and Examples 10, 15 and 37 of Berry, et al. The Examiner further states that while the alkyls on the corresponding R₅ substituent may differ in number, such differences in closely structured related compounds would have been obvious to one of ordinary skill in the art as the resulting products would not have been unexpected.

The Examiner also states that the previously submitted comments regarding the data was not in proper affidavit form and thus not considered.

The Applicants respectfully submit that to establish a *prima facie* case of obviousness, three basic criteria must be met. First, there must be some suggestion or motivation to modify the reference or to combine the reference teachings. Second, there must be a reasonable expectation of success. Third, the prior art reference (or references when combined) must teach or suggest all of the claim limitation. The Applicants

respectfully submit that one skilled in the art would have no suggestion or motivation to modify and/or combine the aforementioned references in order to arrive at the presently claimed invention. Additionally, even if one skilled in the art were to consider the teachings of the cited prior art alone, or in combination, each and every limitation of the present invention would not be disclosed, nor would there be a reasonable expectation of success if the aforementioned references were to be considered either alone or in combination. In addition, prior art must be considered in its entirety, i.e., as a whole (emphasis provided), including portions that would lead away from the claimed invention (M.P.E.P. §2141.02, citing W.L. Gore & Associates, Inc. v. Garlock, Inc., 721 F.2d 1540, 220, USPQ 303 (Fed. Cir. 1983), cert. denied, 469 U.S. 851 (1984)), the proposed modification cannot render the prior art unsatisfactory for its intended purpose or change the principle of operation of a reference (M.P.E.P. §2143.01), and Examiner's conclusion of obviousness may not be based on improper hindsight (M.P.E.P. §2145(X)(A)).

The Applicants respectfully traverse this rejection and respectfully submit that this rejection is in error and should not be maintained. It is first submitted that the prior art fails to teach each and every limitation of the presently amended claims as discussed above.

Accordingly, the rejection should be considered improper.

The Applicants also submit herewith a declaration executed by one of the inventors – Prof. Dr. Heinz Berner – in support of the previously filed comments regarding the data. Those arguments are incorporated herein in their entirety and repeated for the Examiner's ease of reference.

The Applicants had previously enclosed *in vitro* and *in vivo* data (i.e., "screening") of (1) compounds of the present invention (i.e., WO 2004011431, which corresponds to and is the equivalent of the present application), (2) a similar compound of Ascher, et al. '804 and (3) a similar compound of Berry, et al. '128, which are elaborated upon in the attached "Materials and Methods."

In view of the previously attached data, it is respectfully submitted that the data set forth the superiority of the compounds of the present invention when compared with data obtained from compounds of Ascher, et al. '804 and Berry, et al. '128. In particular, it is submitted that the overall results of the data obtained from Examples 3 and 7 of the present application demonstrate excellent *in vitro* activity. Moreover, the aforementioned examples show excellent *in vivo* activities (e.g., $ED_{50} = 8.72$ mg/kg for Example 3 and 16.2 mg/kg for Example 7) as compared to the values from the cited prior art (e.g., $ED_{50} = 8.83$ mg/kg for Ascher, et al. '804 and 26.71 mg/kg for Berry, et al.).

It is further submitted that a comparison of the data of the most prominent example (i.e., retapamulin of Examples 50 and 58) from Berry, et al. with the data of examples of the present invention clearly set forth the improved *in vivo* activity of compounds of Examples 3 and 7 of the present invention. The improved *in vivo* efficacy is also shown by a comparison with the data of the compound of Example 1 of Ascher, et al. '804. It is submitted that these results clearly indicate improved *in vivo* activity of a compound of the present invention.

It is also submitted by the Applicants that one of the most prominent toxicities of pleuromutilins is liver toxicity. The *in vitro* hepatoxicity assay is able to indicate the hepatoxicity potential of the corresponding pleuromutilin derivative. In the *in vitro* hepatoxicity, the

compounds of Example 3 (IC₅₀ = 160 μ g/ml) and Example 7 (IC₅₀ = 131 μ g/ml) of the present invention compared to Example 1 of Ascher, et al. (IC₅₀ = 63 μ g/ml) show remarkable higher IC₅₀ values and therefore have a definitively lower potential for *in vivo* liver toxicity.

In summary, it is submitted that the compounds of Examples 3 and 7 of the present invention have improved *in vivo* activity compared to the compounds of Examples 50 and 58 in Berry, et al. and liver toxicity is decreased compared with the compound of Example 1 of Ascher, et al. Improved efficacy of the compounds of the present invention as shown above could not be expected from either Ascher, et al. or Berry, et al., or even from a combination of the teachings of Ascher, et al. and Berry, et al., and therefore one skilled in the art would lack motivation to refer to and/or modify the teachings of the cited prior art in order to arrive at the presently claimed invention.

In view of the above, withdrawal of this rejection is strongly requested.

Conclusion

In light of the foregoing claims and arguments, it is believed that the present application is in condition for allowance, and such action is earnestly solicited. The Examiner is invited to call the undersigned if there are any remaining issues to be discussed which could expedite the prosecution of the present application.

Respectfully submitted,

Date: april 3, 2009

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Bv:

D. Peter Hochberg Reg. No. 24,603 My name is Prof. Dr. Heinz Berner. I was born on 5.11.194 in Vienna /Austria and my current address is Geyergasse 2a, A-1180 Vienna/Austria.

I declare that,

I am a graduated chemist and professor at the Medicinal Chemistry Department of the University, Althanstrasse 14, A- 1090 Vienna (Austria) and my experience is as follows:

Synthetic Organic Chemistry as well as Synthetic Organic Electrochemistry in the fields of: Steroids and Diterpenes
Steroid Alkaloids (Batrachotoxin)
Amino Acids
Lipopeptides and Cyclopeptolides
Heterocycles

Molecular Modelling

QSAR: Regression-, Principal Component-, Cluster- and Discriminant-Analysis

Curriculum Vitae

Born:

5.11.1940, Vienna (Austria)

High School:

Humanistisches Gymnasium, Vienna, A- 1130, Fichtnergasse 15

Matura 1958

Military Service:

1958/1959

Stüdies:

Chemistry, Universität of Vienna 1959-1967,

Post Doctoral

Fellow:

1968-1970, ETH Zürich: Synthesis of the Steroid Alkaloid

Batrachotoxin.

Habilitation:

In the field of "Pharmaceutical Chemistry"at the Formal-und

Naturwissenschaftliche Fakultät der Universität Wien, 1983.

Further Studies:

Synthetic Organic Electrochemistry, University of Lund, Sweden,

Department Prof. Eberson, April-June 1983.

Industry:

Since 1970 at Novartis Research Institute (Former Sandoz Research Institute, - since 1996 at Novartis/Sandoz/ABRI and successor firm

NABRIVA until end of 2005.

Synthesis and cytotoxic activity of resveratrol-based compounds. Handler, Norbert; Saiko, Philipp; Jaeger, Walter; Szekeres, Thomas; Wacheck, Volker; Berner, Heinz; Leisser, Klaus; Erker, Thomas. Department of Medicinal Chemistry, University of Vienna, Vienna, Austria. Monatshefte fuer Chemie (2008), 139(5), 575-578. Publisher: Springer Wien, CODEN: MOCMB7 ISSN: 0026-9247. Journal written in English. AN 2008:513088 CAPLUS (Copyright (C) 2009 ACS on SciFinder (R))

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Constitution of tormentol (tormentoside). Pailer, Matthias; Berner, Heinz. Univ. Vienna, Vienna, Austria. Monatshefte fuer Chemie (1967), 98(5), 2082-8. CODEN: MOCHAP Journal written in German. CAN 68:22199 AN 1968:22199 CAPLUS (Copyright (C) 2009 ACS on SciFinder (R))

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Already 1970 in Sandoz (later Novartis) I have been involved in pleuromutilin investigations, mainly, but not exclusively, in chemical synthesis of novel pleuromutilins. In 1983 I have found the compound valuemulin which is the active ingredient in Econor®, which is a veterinary antibiotic and I am one of the inventors of the basis valuemulin patent:

Pleuromutilin derivatives and their use. Berner, Heinz; Vyplel, Hermann. (Sandoz A.-G., Switz.; Sandoz-Patent-G.m.b.H.; Sandoz-Erfindungen Verwaltungsgesellschaft m.b.H.). Eur. Pat. Appl. (1985), 16 pp. CODEN: EPXXDW EP 153277 A2 19850828 Designated States R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE. Patent written in German. Application: EP 85-810049 19850211. Priority: DE 84-3405632 19840217; DE 84-3413708 19840412.

Since 1996 in Novartis/Sandoz/ABRI and later in the successor firm NABRIVA novel pleuromutilins have been developed under my supervision.

- I am one of the inventors designated in US 2005/250811, hereinafter designated as "present invention". I have contributed to the present invention by provision of novel pleuromutilins and I have further contributed by involvement in the design and the results of activity and toxicity tests of compounds of the present invention.

I further declare that, to the best of my knowledge

- I have read and understood the inventive step argument raised by the Examiner in the Official Action issued on 05/29/2008 and in the Official Action issued on 01/29/2009.

- I also have read the response filed to the first Official Action on August 29, 2008 and I fully agree with its content.
- The EXPERIMENTS as attached to that Declaration which have been filed also in response to the first Official Action (together with the RESULTS and SUMMARY as indicated below) have been carried out at Nabriva/Novartis/Sandoz/ABRI. In these EXPERIMENTS activity and toxicity of compounds of the present invention, namely such as designated in the EXPERIMENTS, have been compared with activity and toxicity of compounds of the prior art, namely compounds such as designated in the EXPERIMENTS. The obtained results are set out in the table of the RESULTS as attached herewith.

I have considered the RESULTS obtained in these EXPERIMENTS and from the RESULTS I conclude that the compounds of the present invention have outstanding high activity and low toxicity compared with the compounds of the prior art, e.g. as set out in the SUMMARY attached herewith.

-I further conclude that, from the prior art it could not be expected that the compounds of the present invention could show such outstanding high activity and low toxicity compared with those of the prior art compounds, e.g. because of certain similarities in chemical structure of compounds of the present invention compared with the prior art compounds. Thus, I further conclude that the high activity and low toxicity of the compounds of the present invention is unobvious from the prior art and thus inventive.

Vienna/Austria

3rd of March 2009

Signature

Prof Cheinth

Prof. Dr. Heinz Berner